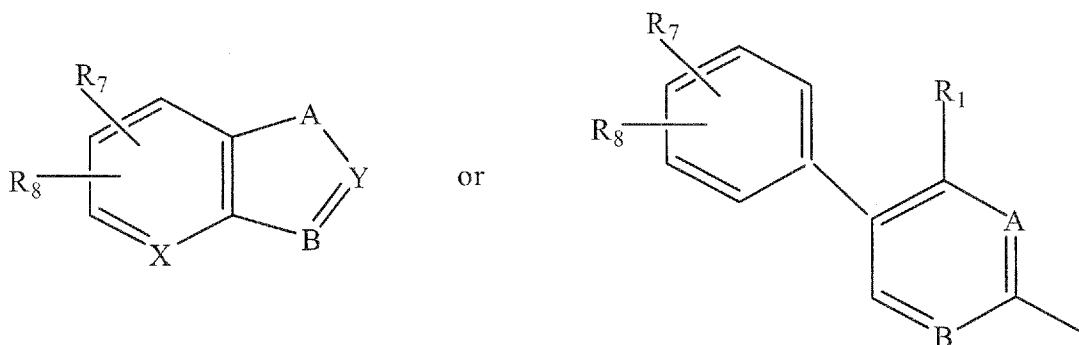


IN THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1-34. (Canceled)

35. (New) A method of inhibiting HIV replication, said method comprising contacting a cell comprising HIV with an effective amount of a compound having the structure:



wherein

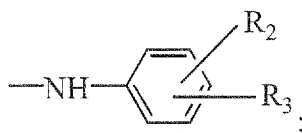
A is selected from the group consisting of N, CR₁, and $\begin{array}{c} \text{R}_1 \\ | \\ \text{---CHN---} \end{array}$;

B is selected from the group consisting of N and S;

Y is selected from the group consisting of Se, CH and CR₄;

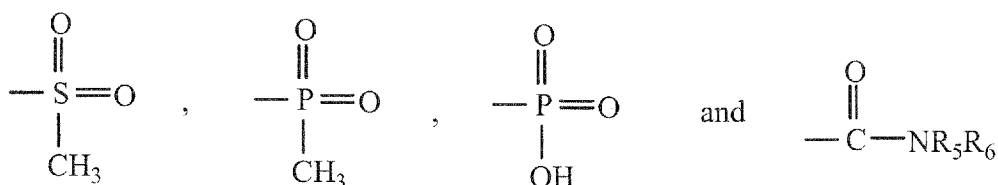
X is selected from the group consisting of CH and N;

R₁ is selected from the group consisting of H, NR₅R₆ and



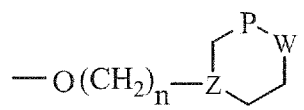
R₂ and R₃ are independently selected from the group consisting of H, halo, hydroxy and C₁-C₄ alkyl;

R₄ is selected from the group consisting of H, halo, hydroxy and C₁-C₄ alkyl;



R₅ and R₆ are independently selected from the group consisting of H and C₁-C₄ alkyl;

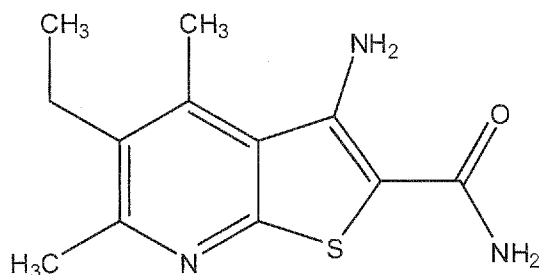
R₇ and R₈ are independently selected from the group consisting of H, halo, hydroxy, C₁-C₄ alkyl, C₁-C₄ alkoxy, -NHC(O)CH₃ and -O(C₁-C₄ alkyl)(C₅-C₆ heterocyclic) or R₇ and R₈ together with the atoms to which they are attached form an optionally substituted C₅-C₆ aryl, wherein the aryl ring is optionally substituted with halo, C₁-C₄ alkyl, C₁-C₄ alkoxy, C₁-C₄ alkyl(C₅-C₆ aryl) and -O(C₁-C₄ alkyl)(C₅-C₆ heterocyclic). In one embodiment Y is CR₄, R₇ is H or C₁-C₄ alkoxy, R₈ is halo or



, wherein n is an integer ranging from 1-5, and P, W and Z are independently selected from the group consisting of O, S, CH₂ and NH;

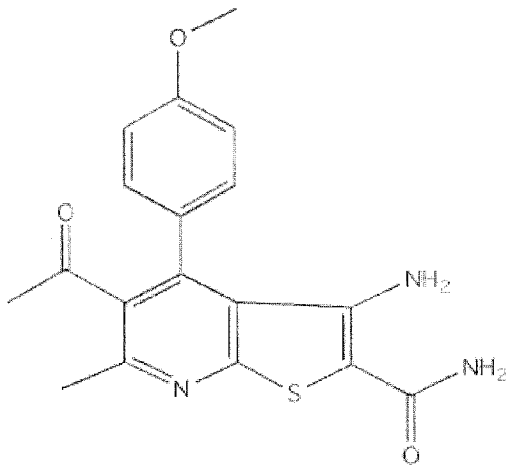
further wherein said compound is selected from the group consisting of

103833:

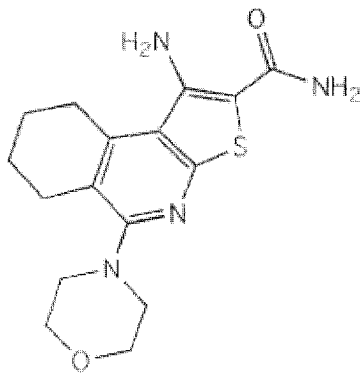


3-amino-5-ethyl-4,6-dimethylthieno[2,3-b]pyridine-2-carboxamide,

AE-641/15124054 (311)

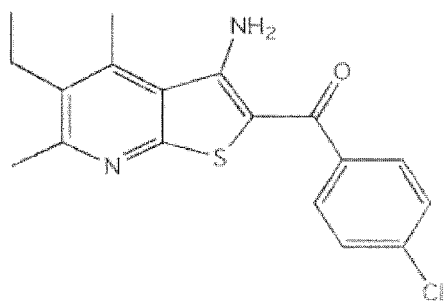


AE-848/11105217 (312)



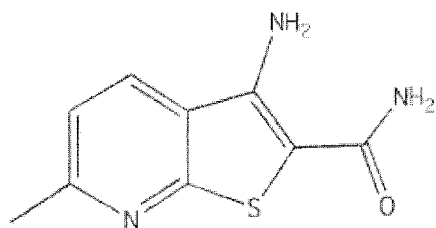
(313)

AE-848/34405027



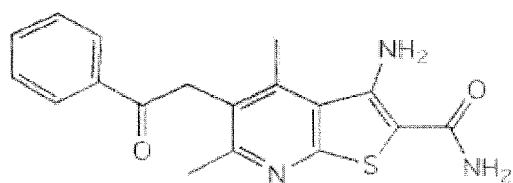
(314)

AG-205/08592044

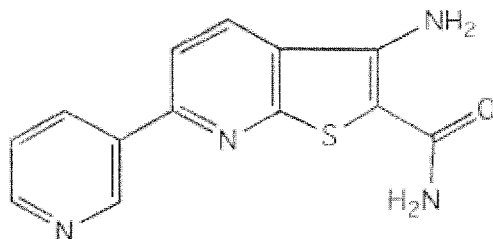


(315)

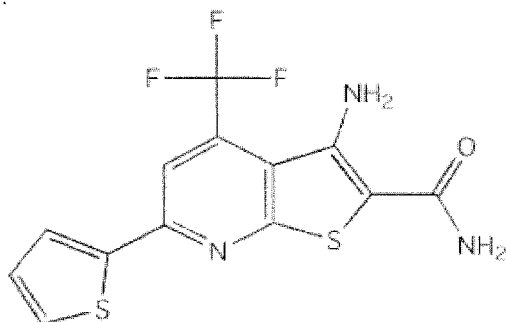
AG-205/11781740



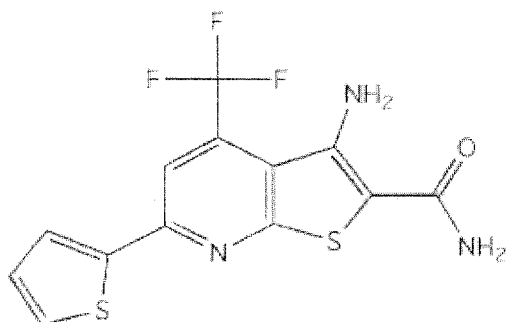
AG-205/31312022 (316)



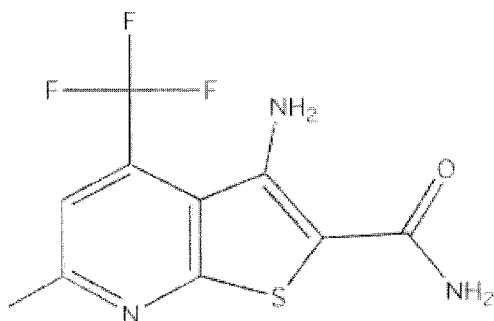
AG-205/33137032 (317)



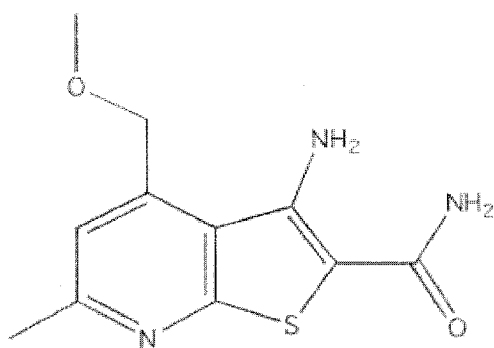
AG-205/33137032 (318)



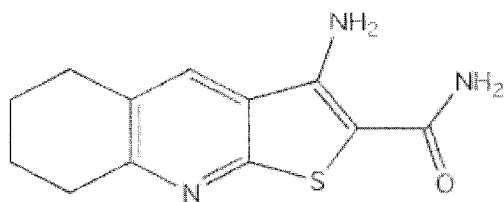
AG-205/33139015 (319)



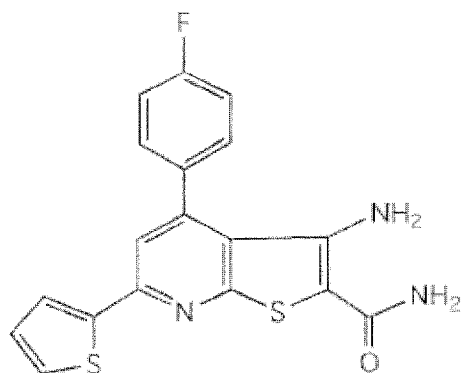
AG-205/33156001 (320)



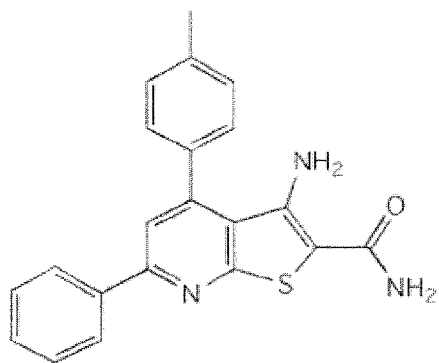
AG-205/33684025 (321)



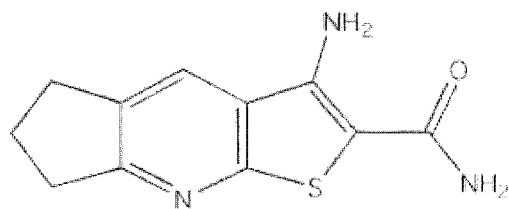
AG-205/36992106 (322)



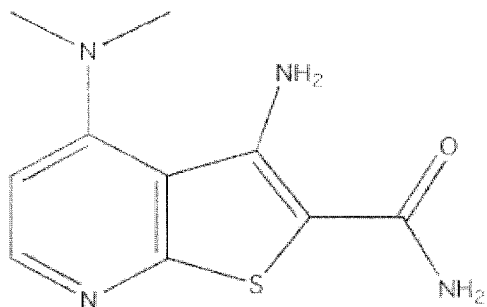
AG-690/1370414 (323)



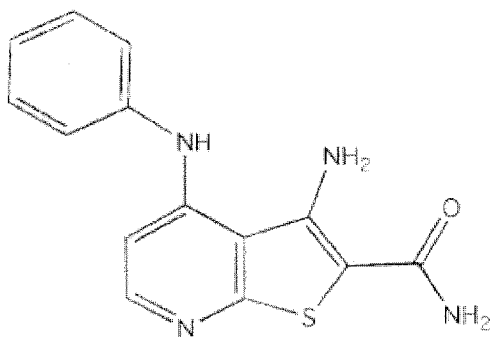
AG-690/34037018 (324)



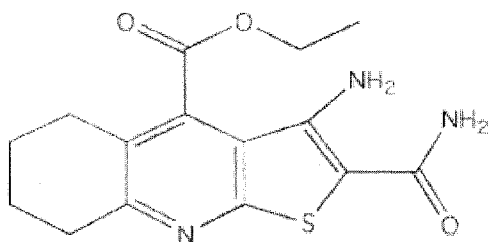
AH-262/343350 (325)



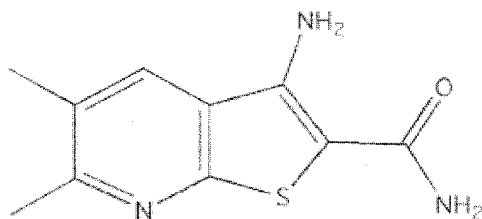
AH-262/36083007 (326)



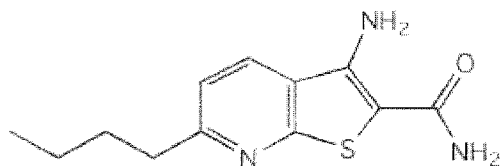
AK-777/115000 (327)



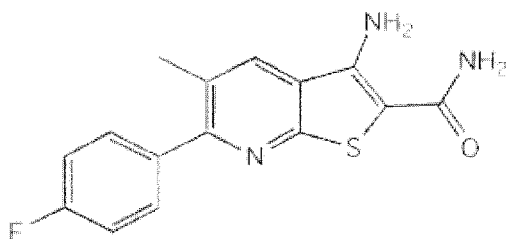
AK-777/36935027 (328)



AK-968/37156085 (329)

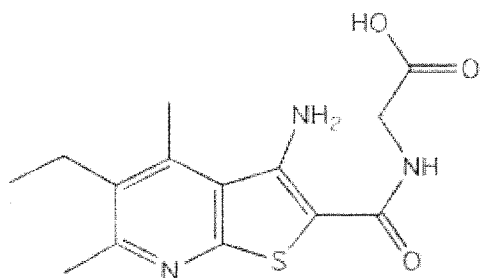


AM-807/12740245 (330)



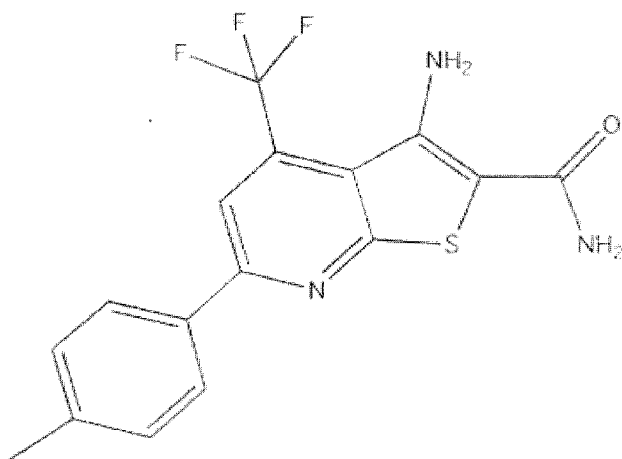
AM-807/13614287

(331)



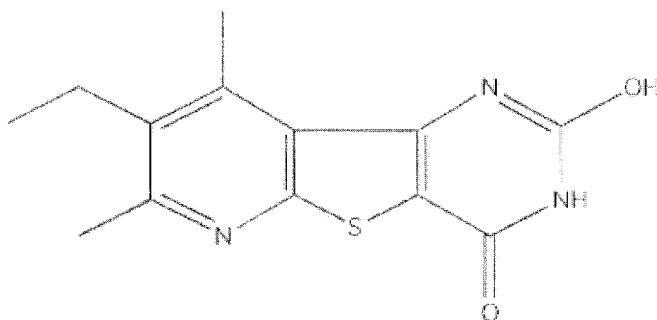
AM-807/14147906

(332)

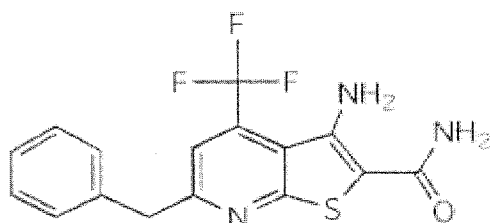


AM-807/42004022

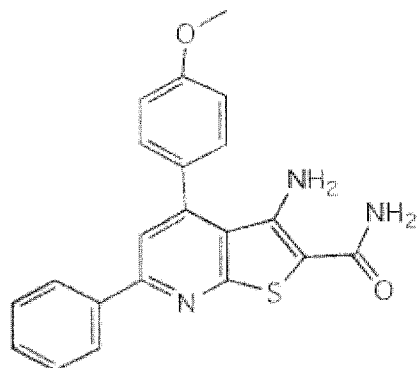
(333)



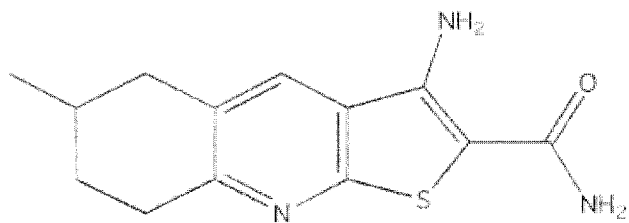
AM-807/42860050 (334)



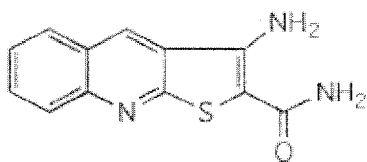
AN-329/05740035 (335)



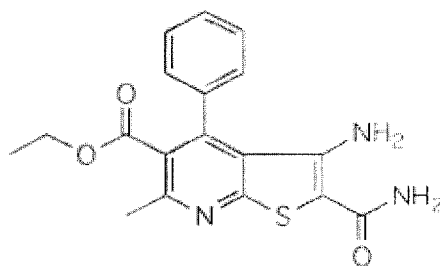
AN-919/14791006 (336)



AO-799/42008042 (337)



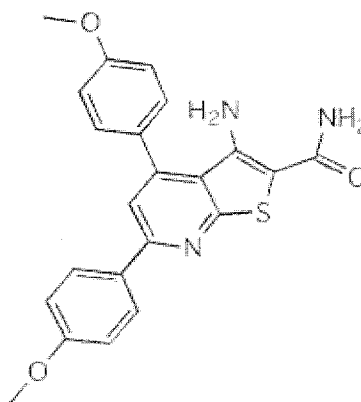
AO-799/43115183 (338)



and

(339)

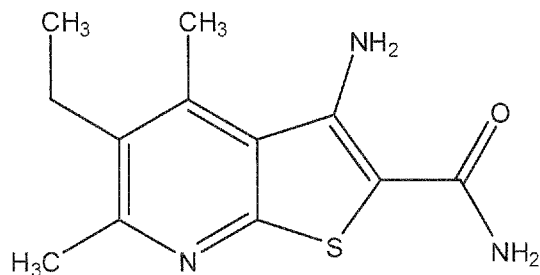
AQ-750/42052143



36. (New) The method of claim 35, wherein said compound inhibits REV function.

37. (New) The method of claim 35, wherein HIV virion production is dependent on Rev protein expression.

38. (New) The method of claim 35, wherein said compound is 103833:



3-amino-5-ethyl-4,6-dimethylthieno[2,3-b]pyridine-2-carboxamide